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PC32213A

PTO/SB/08A (07-05)

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INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
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Complete if Known

Application Number

Filing Date

First Named Inventor Yoshinobu Hashizume, et al.

Art Unit

Examiner Name

Sheet

Attorney Docket Number

U. S. PATENT DOCUMENTS Cite Document Number Publication Date Name of Patentee or Pages, Columns, Lines, Where Initials* MM-DD-YYYY Applicant of Cited Document Relevant Passages or Relevant Number-Kind Code^{2 (F known)} Figures Appear US-US-HS. US-US-US-US-US-US-US-US-US: US-US-US-US-US-

FOREIGN PATENT DOCUMENTS							
Examiner Initials*	Cite No.1	Foreign Patent Document	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages Or Relevent Figures Appear		
		Country Code ³ "Number ⁴ "Kind Code ⁵ (if known)				T ⁸	
		JP2002173485 (Equiv WO 2002/026714 A1)	06-21-2002	Takeda Chem Ind Ltd		\checkmark	
		WO 1994/029309 A1	12-22-1994	Merck & Co Inc			
		WO 1998/025605 A1	12-12-1997	Merck & Co Inc			
		WO 2002/085354 A1	10-31-2002	Euroceltique SA			
		WO 2003/000677 A1	01-03-2003	Pfizer Pharmaceuticals	Inc		
		WO 2003/064425 A1	08-07-2003	Pfizer Pharmaceuticals	Inc		

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STATEMENT BY APPLICANT			PPLICANT	First Named Inventor	Yoshinobu Hashizume, et al.		
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Sheet	2	of	2	Attorney Docket Number	PC32213A		

Examiner	Cite	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of	
nitials*	Cite No. ¹	the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
		CALO, G. et al., "Pharmacology of Nociceptin and its Receptor: a Novel Therapeutic Target", British Journal of Pharmacology, 2000, pp 1261-1283, Vol 129	
-		MEUNIER, J. et al., "Isolation and Structure of the Endogenous Agonist of Opioid Receptor-like ORK1 Receptor", Neture, 1995, pp 532-535, Voi 377	
		REINSHEID, R. K. et al., "Orphanin FQ: A Neuropeptide That Activates an Opioidlike G Protein-Coupled Receptor", Science, 1995, pp 792-794, Vol 270, No 5237	
		RONZONI, S. et al., "Lead Generation and Lead Optimisation Approaches in the Discovery of Selective Non-peptide ORL-1 Receptor Agonists and Antagonists", Expert Opinion on Therapeutic Patents, 2001, pp 525-546, Vol 11, No 4	
		SCHMIDIT, A. W. et al., "The Novel Antipsychotic Ziprasidone has e Unique Human Receptor Binding Profile Compered to Other Agents", Society for Neuroscience, 1998, pp 2177, Vol 24	
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